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//046237 A

(54) Title: HUMANIZED ANTIBODIES THAT RECOGNIZE BETA AMYLOID PEPTIDE

(57) Abstract: The invention provides improved agents and methods for treatment of diseases associated with amyloid deposits of Aß in the brain of a patient. Preferred agents include humanized antibodies.

Internation Application No PCT/US 01/46587

A. CLASSIFICATION OF SUBJECT MATTER IPC 7 C07K16/18 A61P25/28 C12N15/13 C12N15/85 A61K39/395 C12N5/10

According to International Patent Classification (IPC) or to both national classification and IPC

B. FIELDS SEARCHED

Minimum documentation searched (classification system followed by classification symbols)

IPC 7 C07K

Documentation searched other than minimum documentation to the extent that such documents are included in the fields searched

Electronic data base consulted during the international search (name of data base and, where practical, search terms used)

CHEM ABS Data, SEQUENCE SEARCH, MEDLINE, BIOSIS, EPO-Internal, WPI Data, PAJ

C. DOCUM	ENTS CONSIDERED TO BE RELEVANT		· · · · · · · · · · · · · · · · · · ·
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χ Fur	ther documents are listed in the continuation of box C.	χ Patent family members are listed	in annex.
"A" docum consi "E" earlier filing "L" docum which citatic "O" docum other "P" docum later	ategories of cited documents: and defining the general state of the art which is not dered to be of particular relevance document but published on or after the international date ent which may throw doubts on priority claim(s) or a is cited to establish the publication date of another or or other special reason (as specified) then treferring to an oral disclosure, use, exhibition or means lent published prior to the international filing date but than the priority date claimed	"T" later document published after the intro or priority date and not in conflict with cited to understand the principle or the invention." X" document of particular relevance; the cannot be considered novel or cannot involve an inventive step when the document of particular relevance; the cannot be considered to involve an indocument is combined with one or ments, such combination being obvious in the art. "&" document member of the same patent Date of mailing of the international se	the application but eory underlying the claimed invention to be considered to bournent is taken alone claimed invention enter such docurrent to a person skilled
2	April 2003	17.04.2003	
Name and	mailing address of the ISA European Patent Office, P.B. 5818 Patentlaan 2 NL – 2280 HV Rijswijk Tel. (+31–70) 340–2040, Tx. 31 651 epo nl, Fax: (+31–70) 340–3016	Authorized officer COVONE-VAN HEES,	

Internation Application No
PCT/US 01/46587

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	Relevant to claim No.
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Internation Application No
PCT/US 01/46587

Citation of document, with indication, where appropriate, of the relevant passages Relevant to claim No. Relevant to claim No. Residuate the Monocolonal Antibody By CDR-GRAFTING: THE IMPORTANCE OF FRAMEWORK RESIDUES ON LOOP CONFORMATION" PROTEIN ENGINEERING, OXFORD UNIVERSITY PRESS, SURREY, GB, vol. 4, no. 7, 1 October 1991 (1991–10–01), pages 773–783, XP002048549 ISSN: 0269–2139 the whole document	o
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Intertarional application No. PCT/US 01/46587

Box I	Observations where certain claims were found unsearchable (Continuation of item 1 of first sheet)
This Inte	ernational Search Report has not been established in respect of certain claims under Article 17(2)(a) for the following reasons:
1. χ	Claims Nos.: because they relate to subject matter not required to be searched by this Authority, namely:
	Although claims 58-61,133-136 are directed to a method of treatment of the human/animal body, the search has been carried out and based on the alleged effects of the compound/composition.
2. X	Claims Nos.: 69,70,144,145 because they relate to parts of the International Application that do not comply with the prescribed requirements to such an extent that no meaningful International Search can be carried out, specifically:
	see FURTHER INFORMATION sheet PCT/ISA/210
з. 🗌	Claims Nos.: because they are dependent claims and are not drafted in accordance with the second and third sentences of Rule 6.4(a).
Box II	Observations where unity of invention is lacking (Continuation of item 2 of first sheet)
This Inte	ernational Searching Authority found multiple inventions in this international application, as follows:
	see additional sheet
1. X	As all required additional search fees were timely paid by the applicant, this International Search Report covers all searchable claims.
2.	As all searchable claims could be searched without effort justifying an additional fee, this Authority did not invite payment of any additional fee.
з	As only some of the required additional search fees were timely paid by the applicant, this International Search Report covers only those claims for which fees were paid, specifically claims Nos.:
4.	No required additional search fees were timely paid by the applicant. Consequently, this International Search Report is restricted to the invention first mentioned in the claims; it is covered by claims Nos.:
Remark	on Protest The additional search fees were accompanied by the applicant's protest.
	χ No protest accompanied the payment of additional search fees.

FURTHER INFORMATION CONTINUED FROM PCT/ISA/ 210

This International Searching Authority found multiple (groups of) inventions in this international application, as follows:

1. Claims: 1-83,138-141,155,156 (complete) 149 (partially)

Humanized immunoglobulin (Ig) light chain and heavy chain comprising (i) variable region complementarity determining regions (CDRs) from 3D6 Ig light and heavy chain variable regions sequence set forth as seq ID. 2 and 4 respectively, and (ii) variable framework region from a human acceptor Iq light and heavy chain, provided that at least one framework residue is substituted with the corresponding amino acid residue from the mouse 3D6 light and heavy chain variable region sequence. Furthermore, the subject matter comprises the related products and methods: chimeric Ig comprising CDRs from 3D6 Ig or antigen binding fragment thereof, comprising a variable heavy and light chain (set forth as seq. ID 8, 12 and seq. ID 5,11 respectively), humanized antibody, method of preventing or treating an amyloidogenic disease and Alzheimer in patients, pharmaceutical compositions comprising said Ig, isolated polypeptide or fragments (as referred to in claims 63-66), a variant of said polypeptide, nucleic acid, vectors and host cells to encode and produce said Ig, method to produce said Ig. Method for identifying residues amenable to substitution in a humanized 3D6 variable framework region and use of the variable sequence to produce a three dimensional image of 3D6 Ig.

2. Claims: 84-137,142-148,150-154,157, 158 (complete) 149 (partially)

> Humanized immunoglobulin (Ig) light chain and heavy chain comprising (i) variable region complementarity determining regions (CDRs) from 10D5 Ig light and heavy chain variable regions sequence set forth as seq ID. 14 and 16 respectively, and (ii) variable framework region from a human acceptor Ig light and heavy chain, provided that at least one framework residue is substituted with the corresponding amino acid residue from the mouse 10D5 light and heavy chain variable region sequence. Furthermore, the subject matter comprises the related products and methods: chimeric Ig, humanized antibody, method of preventing or treating an amyloidogenic disease and Alzheimer in patients, pharmaceutical compositions comprising said Ig, isolated polypeptide, a variant of said polypeptide, nucleic acid, vectors and host cells to encode and produce said Ig, method to produce said Ig. Method for identifying residues amenable to substitution in a humanized 10D5 variable framework region and use of the variable sequence to produce a three dimensional image of 10D5 Ig.

FURTHER INFORMATION CONTINUED FROM PCT/ISA/ 210

Continuation of Box I.2

Claims Nos.: 69,70,144,145

Present claims 69,70,144,145 relate to an extremely large number of possible products. In fact, the claims contain so many possible variants that a lack of clarity (and/or conciseness) within the meaning of Article 6 PCT arises to such an extent as to render a meaningful search of the claims impossible. Consequently, the search has not been carried out.

The applicant's attention is drawn to the fact that claims, or parts of claims, relating to inventions in respect of which no international search report has been established need not be the subject of an international preliminary examination (Rule 66.1(e) PCT). The applicant is advised that the EPO policy when acting as an International Preliminary Examining Authority is normally not to carry out a preliminary examination on matter which has not been searched. This is the case irrespective of whether or not the claims are amended following receipt of the search report or during any Chapter II procedure.

intermation on patent ramily members

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PCT/US 01/46587

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